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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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10/766,403

01/27/2004

Luiz Belardinelli

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EXAMINER

CRANE, LAWRENCE E

ART UNIT

PAPER NUMBER

1623

SHORTENED STATUTORY PERIOD OF RESPONSE	MAIL DATE	DELIVERY MODE
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3 MONTHS

04/10/2007

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

Office Action Summary	Application No. 10/766,403	Applicant(s) BELARDINELLI ET AL.	
	Examiner L. E. Crane	Art Unit 1623	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 25 August 2006.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1,5-8,10-17,19-31,62 and 63 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1,5-8,10-17,19-31,62 and 63 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☒ The drawing(s) filed on 16 August 2004 is/are: a) ☒ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)
Paper No(s)/Mail Date <u>12/21/2006</u> . | 6) <input type="checkbox"/> Other: _____ |

This application has been filed with informal drawings which are acceptable for examination purposes only. Formal drawings will be required when the application is allowed.

Claims **2-4, 9, 18 and 32-61** have been cancelled, claims **1, 5, 8, 10-11, 16, 19, 21, 23, 25 and 28** have been amended, the abstract of the disclosure has been amended, and new claims **62 and 63** have been added as per the amendment filed August 26, 2006. One additional Information Disclosure Statement (1 IDS) filed December 21, 2006 has been received with all cited references and made of record.

Claims **1, 5-8, 10-17, 19-31 and 62-63** remain in the case.

Note to applicant: when a rejection refers to a claim **X** at line **y**, the line number “**y**” is determined from the claim as previously submitted by applicant in the most recent response including ~~lines deleted by line through~~.

Claim **1** is rejected under 35 U.S.C. §112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

In claim **1** at line 11, the term “co-solvent comprises methylboronic acid or borate buffer” is technically inaccurate because “methylboronic acid” is a solid at room temperature (m.p. 91-94 degrees C). Appropriate amendment to reflect the probable role of methylboronic acid as an optional -- solute -- is respectfully requested. See also claim **5** wherein the same error occurs.

Applicant's arguments with respect to claim **1** have been considered but are moot in view of the new grounds of rejection.

The non-statutory double patenting rejection, whether of the obviousness-type or non-obviousness-type, is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the “right to exclude” granted by a patent. *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); *In re Van Ornam*, 686 F. 2d 937, 214 USPQ 761 (CCPA 1982); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir 1985); and *In re Goodman*, 29 USPQ 2d 2010 (Fed. Cir. 1993).

A timely filed terminal disclaimer in compliance with 37 C.F.R. § 1.321(b) and (c) may be used to overcome an actual or provisional rejection based on a non-statutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 C.F.R. §1.78(d).

Effective January 1, 1994, a registered attorney or agent or record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 C.F.R. §3.73(b).

Claims 1, 5-8, 10-17, 19-31 and 62-63 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-11 of copending Application No. 11/253,322. Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of imaging and the alleged active ingredient (CVT-3164) are directed to substantially overlapping subject matter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant has noted the above ground of rejection but has not supplied a Terminal Disclaimer or otherwise responded. Therefore the above ground of rejection has been maintained.

Claims 1, 5-8, 10-17, 19-31 and 62-63 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-30 of copending Application No. 10/629,368. Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment and the alleged active ingredients are directed to substantially overlapping subject matter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant has noted the above ground of rejection but has not supplied a Terminal Disclaimer or otherwise responded. Therefore the above ground of rejection has been maintained.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **11, 14-27, 29-30, 34 and 36-37** of copending Application No. **11/070,768**. Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment and the alleged active ingredients are directed to substantially overlapping subject matter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant has noted the above ground of rejection but has not supplied a Terminal Disclaimer or otherwise responded. Therefore the above ground of rejection has been maintained.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **2-4** of U. S. Patent No. **7,109,180** (PTO-892 ref. A). Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment wherein either coronary vasodilation, or increased coronary blood flow made possible by said vasodilation, is induced by administration of the identical active ingredient, CVT-3033. Therefore the two applications are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection. Applicant's newly cited prior art necessitated this new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **5-8 and 10-22** of U. S. Patent **7,183,264** (PTO-892 ref. **B**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the claimed methods of treatment in both applications involve administration of the identical active ingredient, CVT-3146, to induce coronary vasodilation for the purpose of cardiac blood flow imaging. Therefore, the instant claims sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection. Applicant's newly cited prior art necessitated this new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **10-24** of U. S. Patent No. **7,144,872** (PTO-1449 (#5) ref. **E5**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the methods of treatment in both applications involve administration of the identical active ingredient, CVT-3146, to induce coronary vasodilation for the purpose of cardiac blood flow imaging. Therefore, the instant claims sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection. Applicant's newly cited prior art necessitated this new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **9, 10 and 16** of U. S. Patent No. **6,641,210** (PTO-1449 (#3) ref. **A15**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the methods of treatment in both the application and the patent involve administration of the identical active ingredient, CVT-3146, to induce coronary vasodilation for the purpose of cardiac blood flow imaging. Therefore, the instant claims sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **2-4** of U. S. Patent No. **6,770,634** (PTO-1449 (#3) ref. **A17**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment wherein either coronary vasodilation, or increased coronary blood flow made possible by said vasodilation, is induced by administration of the identical active ingredient, CVT-3033. Therefore the two claim sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **29-31** of U. S. Patent No. **6,214,807** (PTO-1449 (#1) ref. **A12**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the methods of treatment in both the application and the patent involve administration of the identical active ingredients, CVT-3033 or CVT-3146, to induce coronary vasodilation for the purpose of cardiac blood flow imaging. Therefore, the instant claims sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **11-13** of U. S. Patent No. **6,403,567** (PTO-1449 (#1) ref. **A13**). Although the conflicting claims are not identical, they are not patentably distinct from each other because the methods of treatment in both the application and the patent involve administration of the identical active ingredient, CVT-3146, to induce coronary vasodilation for the purpose of cardiac blood flow imaging. Therefore, the instant claims sets are directed to substantially overlapping subject matter.

Applicant's arguments with respect to claims **1-61** have been considered but are deemed to be moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims **2-4** of U. S.

Application No. 11/588,834 (PTO-1449 (#5) ref. D5). Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment wherein either coronary vasodilation, or increased coronary blood flow made possible by said vasodilation, is induced by administration of the identical active ingredient, CVT-3033 . Therefore the two applications are directed to substantially overlapping subject matter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicant's arguments with respect to claims 1-61 have been considered but are deemed to be moot in view of the new grounds of rejection. Applicant's newly cited prior art necessitated this new grounds of rejection.

Claims 1, 5-8, 10-17, 19-31 and 62-63 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 29-31 of U. S. Application No. 11/522,120 (PTO-1449 (#5) ref. C5). Although the conflicting claims are not identical, they are not patentably distinct from each other because the method of treatment wherein either coronary vasodilation, or increased coronary blood flow made possible by said vasodilation, is induced by administration of the identical active ingredient, CVT-3146. Therefore the two applications are directed to substantially overlapping subject matter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicant's arguments with respect to claims 1-61 have been considered but are deemed to be moot in view of the new grounds of rejection. Applicant's newly cited prior art necessitated this new grounds of rejection.

The following is a quotation of the appropriate paragraphs of 35 U.S.C. §102 that form the basis for the rejections under this section made in this Office action:

"A person shall be entitled to a patent unless -

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent."

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States."

(e) the invention was described in

(1) an application for patent described under section 122(b), by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effect under this subsection of a national application filed under this subsection of a national application published under section 122(b) only if the international application designating the United States was published under Article 21(2)(a) of such treaty in the English language; or

(2) a patent granted on an application by another filed in the United States before the invention by the applicant for patent, except that a patent shall not be deemed filed in the United States for the purposes of this subsection based on the filing of an international application filed under the treaty defined in section 351(a)."

(f) he did not himself invent the subject matter sought to be patented."

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(f) is claimed in part or identically by a different inventive entity identified in US Patent Application **11/522,120** (PTO-1449 (#5) ref. C5).

Applicant is respectfully requested to resolve the differences in inventorship or to take other appropriate action.

Applicant's arguments with respect to claims **1, 5-8, 10-17 and 19-31** have been considered but are moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(f) is claimed in part or identically by a different inventive entity identified in US Patent Application **11/588,834** (PTO-1449 (#5) ref. D5).

Applicant is respectfully requested to resolve the differences in inventorship or to take other appropriate action.

Applicant's arguments with respect to claims **1, 5-8, 10-17 and 19-31** have been considered but are moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(f) is claimed in part or identically by a different inventive entity identified in US Patent Application **10/629,638** (PTO-1449 (#5) ref. A5).

Applicant is respectfully requested to resolve the differences in inventorship or to take other appropriate action.

Applicant's arguments with respect to claims **1, 5-8, 10-17 and 19-31** have been considered but are moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(f) is claimed in part or identically by a different inventive entity identified in US Patent Application **11/253,322** (PTO-1449 (#5) ref. **B5**).

Applicant is respectfully requested to resolve the differences in inventorship or to take other appropriate action.

Applicant's arguments with respect to claims **1, 5-8, 10-17 and 19-31** have been considered but are moot in view of the new grounds of rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(b) as being anticipated by **Zablocki et al. '807** (PTO-1449 ref. **A12**).

Applicant is referred to claims **1 and 29-33** wherein adenosine agonists with A_{2A} receptor selectivity are disclosed as part of a pharmaceutical composition and the utility in inducing localized vasodilation for cardiac imaging purposes are disclosed. The compound also known as CVT-3033 may be found at column 20, lines 40-50.

The **CV Therapeutics '778** reference (PTO-1449 ref. **B3**) is the PCT equivalent to the above reference and anticipates for the same reasons.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant argues that the details of the claims newly introduced by amendment distinguish the instant claims over the cited prior art claims. Examiner respectfully disagrees, and notes that method claim **29** in the cited prior art is generic to "imaging of the heart" following administration of one of the compounds now specified herein and therefore inherently encompasses all of the necessary details one of ordinary skill would have deduced in the course of routine experimentation for the purpose of optimizing the prior art. In essence, because the cited prior art is commonly assigned, applicant's assignee already owns the details newly added to the instant claims.

For this reasons the instant rejection is deemed to remain valid and has been maintained.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(e) as being anticipated by **Zablocki et al. '567** (PTO-1449 ref. **A13**).

Applicant is referred to claims **1, 8, 10 and 11-13** wherein the compound, also known as CVT-3164, is disclosed as part of a pharmaceutical composition and as having utility in the imaging of mammalian cardiac circulatory systems.

See also **CV Therapeutics '779** (PTO-1449 ref. **B2**) which is the PCT equivalent to the '**567** reference and also anticipates the instant noted claims for the same reasons.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant is referred to the response to the immediately preceding rejection.

Claims **1, 5-8, 10-17, 19-31 and 62-63** are rejected under 35 U.S.C. §102(a) and/or (b) as being anticipated by **Gao et al.** (PTO-1449 ref. **C2**).

Applicant is referred to the reference at its abstract wherein both CVT-3033 and CVT-3164 are disclosed as having the desirable properties of inducing short term coronary vasodilation during myocardial imaging in the presence of radionuclides. The copy of the reference indicates a publication date of July, 2001, which without more complete date information is deemed to be sufficient to render the instant claimed subject matter anticipated.

Applicant's arguments filed August 25, 2006 have been fully considered but they are not persuasive.

Applicant argues that the lack of identical language means that **Gao et al.** does not anticipate the instant claims. Examiner respectfully disagrees. Applicant is referred to **Gao et al.** at page 215 at column 1, beginning in the "Discussion" section: in particular the second sentence of the first paragraph of this section refers specifically to CVT-3164 and that both this compound and a related compound (CVT-3033) induced "coronary vasodilation" but that "the duration of their effect was remarkably shorter." This disclosure anticipates instant claim **21** wherein "coronary vasodilation" is asserted as the primary effect of CVT-3164. The

associated effects are deemed to be inherently a consequence of "coronary vasodilation." "Coronary vasodilation" is a phenomenon induced by administration of CVT-3164 and is a variable controllable by routine variation of dosage, a variable clearly within the purview of the ordinary practitioner seeking to optimize the prior art via routine experimentation. Therefore, the details of the noted effects not presenting, or presenting, respectively, are deemed to be limitations that do not permit applicant to avoid the finding of anticipation.

The following is a quotation of 35 U.S.C. §103(a) which forms the basis for all obviousness rejections set forth in this Office action:

"A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made."

The above Office action contains numerous new grounds of rejection and therefore could not be made final.

Papers related to this application may be submitted to Group 1600 via facsimile transmission (FAX). The transmission of such papers must conform with the notice published in the Official Gazette (1096 OG 30, November 15, 1989). The telephone number to FAX (unofficially) directly to Examiner's computer is 571-273-0651. The telephone number for sending an Official FAX to the PTO is 571-273-8300.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Examiner L. E. Crane whose telephone number is **571-272-0651**. The examiner can normally be reached between 9:30 AM and 5:00 PM, Monday through Friday.

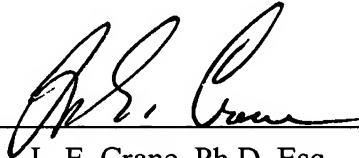
If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ms. S. Anna Jiang, can be reached at **571-272-0627**.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group 1600 receptionist whose telephone number is **571-272-1600**.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published

applications may be obtained from either Private PAIR or Public PAIR. Status Information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see < <http://pair-direct.uspto.gov> >. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

LECrane:lec
03/30/2007



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